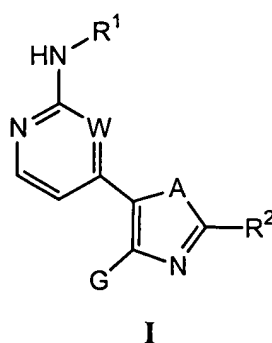


AMENDMENTS TO THE CLAIMS

The following **Listing of Claims** will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Currently amended) A compound of formula I:



or a pharmaceutically acceptable derivative thereof, wherein:

W is nitrogen or CH;

G is hydrogen or C₁₋₃ aliphatic wherein one methylene unit of G is optionally replaced by -C(O)-, -C(O)O-, -C(O)NH-, -SO₂-, or -SO₂NH-;

A is -N-T_(n)-R, oxygen, or sulfur;

R¹ is selected from -T_(n)-R or -T_(n)-Ar¹;

each n is independently 0 or 1;

T is a C₁₋₄ alkylidene chain wherein one methylene unit of T is optionally replaced by -C(O)-, -C(O)O-, -C(O)NH-, -SO₂-, or -SO₂NH-;

Ar¹ is a 3-7 membered monocyclic saturated, partially saturated or aromatic ring having 0-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur, or a 8-10 membered bicyclic saturated, partially saturated or aromatic ring having 0-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur, wherein each member of Ar¹ is optionally substituted with one -Z-R³ and one to three additional groups independently selected from -R, halogen, oxo, -NO₂, -CN, -OR, -SR, -N(R)₂, -NRC(O)R, -NRC(O)N(R)₂, -NRCO₂R,

-C(O)R, -CO₂R, -OC(O)R, -C(O)N(R)₂, -OC(O)N(R)₂, -S(O)R, -SO₂R, -SO₂N(R)₂,
-NRSO₂R, -NRSO₂N(R)₂, -C(O)C(O)R, or -C(O)CH₂C(O)R;

each R is independently selected from hydrogen or a C₁₋₆ aliphatic, wherein said aliphatic is optionally substituted with one to three groups independently selected from oxo, -CO₂R', -OR', -N(R')₂, -SR', -NO₂, -NR'C(O)R', -NR'C(O)N(R')₂, -NR'CO₂R', -C(O)R', -OC(O)R', -C(O)N(R')₂, -OC(O)N(R')₂, -S(O)R', -SO₂R', -SO₂N(R')₂, -NR'SO₂R', -NR'SO₂N(R')₂, -C(O)C(O)R', -C(O)CH₂C(O)R', halogen, or -CN, or two R bound to the same nitrogen atom are taken together with that nitrogen atom to form a five or six membered heterocyclic or heteroaryl ring having one to two additional heteroatoms independently selected from oxygen, nitrogen, or sulfur;

each R' is independently selected from hydrogen or C₁₋₆ aliphatic, wherein said aliphatic is optionally substituted with one to three groups independently selected from oxo, -CO₂H, -OH, -NH₂, -SH, -NO₂, -NHC(O)H, -NHC(O)NH₂, -NHCO₂H, -C(O)H, -OC(O)H, -C(O)NH₂, -OC(O)NH₂, -S(O)H, -SO₂H, -SO₂NH₂, -NHSO₂H, -NHSO₂NH₂, -C(O)C(O)H, -C(O)CH₂C(O)H, halogen, or -CN, or two R' bound to the same nitrogen atom are taken together with that nitrogen atom to form a five or six membered heterocyclic or heteroaryl ring optionally having one or two additional heteroatoms independently selected from nitrogen, oxygen, or sulfur;

Z is a C₁-C₆ alkylidene chain wherein up to two non-adjacent methylene units of Z are optionally replaced by -C(O)-, -C(O)O-, -C(O)C(O)-, -C(O)N(R)-, -OC(O)N(R)-, -N(R)N(R)-, -N(R)N(R)C(O)-, -N(R)C(O)-, -N(R)C(O)O-, -N(R)C(O)N(R)-, -S(O)-, -SO₂-, -N(R)SO₂-, -SO₂N(R)-, -N(R)SO₂N(R)-, -O-, -S-, or -N(R)-;

R² is -Q_(n)-Ar²;

Ar² is selected from a 3-7 membered monocyclic saturated, saturated or aromatic ring having 0-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur, or a 8-10 membered bicyclic saturated, saturated or aromatic ring having 0-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur, wherein each member of Ar² is optionally substituted with 1-5 groups independently selected from -Z-R³, -R, halogen, oxo, -NO₂, -CN, -OR, -SR, -N(R)₂, -NRC(O)R, -NRC(O)N(R)₂, -NRCO₂R, -C(O)R, -CO₂R, -OC(O)R, -C(O)N(R)₂, -OC(O)N(R)₂, -S(O)R, -SO₂R, -SO₂N(R)₂, -N(R)SO₂R, -N(R)SO₂N(R)₂, -C(O)C(O)R, or -C(O)CH₂C(O)R;

Q is a C₁₋₃ alkylidene chain wherein up to two non-adjacent methylene units of Q are optionally replaced by -C(O)-, -C(O)O-, -C(O)C(O)-, -C(O)N(R)-, -OC(O)N(R)-, -N(R)N(R)-, -N(R)N(R)C(O)-, -N(R)C(O)-, -N(R)C(O)O-, -N(R)C(O)N(R)-, -S(O)-, -SO₂-, -N(R)SO₂-, -SO₂N(R)-, -N(R)SO₂N(R)-, -O-, -S-, or -N(R)-;

R³ is selected from -Ar³, -R, halogen, -NO₂, -CN, -OR, -SR, -N(R)₂, -NRC(O)R, -NRC(O)N(R)₂, -NRCO₂R, -C(O)R, -CO₂R, -OC(O)R, -C(O)N(R)₂, -OC(O)N(R)₂, -SOR, -SO₂R, -SO₂N(R)₂, -NRSO₂R, -NRSO₂N(R)₂, -C(O)C(O)R, or -C(O)CH₂C(O)R; and

Ar³ is a 5-6 membered saturated, partially saturated, or aromatic ring having 0-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur, wherein each member of Ar³ is optionally substituted with halogen, oxo, -CN, -NO₂, -R', -OR', -N(R')₂, -N(R')C(O)R', -N(R')C(O)N(R')₂, -N(R')CO₂R', -C(O)R', -CO₂R', -OC(O)R', -C(O)N(R')₂, -OC(O)N(R')₂, or -SO₂R';

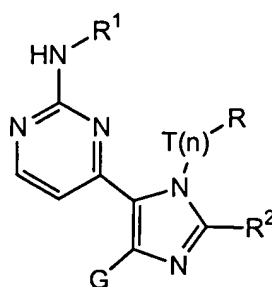
provided that when W is nitrogen and:

(i) A is -N-T_(n)-R and R² is a saturated ring or

(ii) A is sulfur,

then R¹ is other than an optionally substituted phenyl.

2. (Original) The compound according to claim 1, wherein said compound has formula **IIa**:



IIa

or a pharmaceutically acceptable derivative thereof.

3. (Original) The compound according to claim 2, wherein said compound has one or more features selected from the group consisting of:

(a) R¹ is hydrogen, Ar¹ or -T-Ar¹ wherein T is a C₁₋₄ alkylidene chain and Ar¹ is a 6-membered saturated, partially saturated, or aryl ring having zero to two

heteroatoms independently selected from nitrogen, oxygen, or sulfur, and wherein each member of R^1 is optionally substituted with one $-Z-R^3$ and one to three additional groups independently selected from $-CO_2R$, $-OR$, halogen, $-NRSO_2R$, $-SO_2N(R)_2$, $-NRCON(R)_2$, $-NO_2$, or $-N(R)_2$;

- (b) R^2 is Ar^2 or $-CH_2-Ar^2$ wherein Ar^2 is selected from 5-6 membered ring selected from carbocyclic, aryl, or a heterocyclyl or heteroaryl ring having one to two heteroatoms independently selected from nitrogen, oxygen or sulfur, and wherein Ar^2 is optionally substituted with one to five groups independently selected from $-Z-R^3$, $-R$, halogen, $-NO_2$, $-CN$, $-OR$, $-SR$, $-N(R)_2$, $-NRC(O)R$, $-NRC(O)N(R)_2$, $-NRCO_2R$, $-C(O)R$, $-CO_2R$, $-C(O)N(R)_2$, $-OC(O)N(R)_2$, $-S(O)R$, $-SO_2R$, $-SO_2N(R)_2$, $-N(R)SO_2R$, $-N(R)SO_2N(R)_2$, $-C(O)C(O)R$, or $-C(O)CH_2C(O)R$; and
- (c) G is hydrogen.

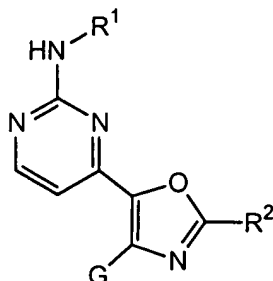
4. (Original) The compound according to claim 3, wherein said compound has one or more features selected from the group consisting of:

- (a) R^1 is selected from a phenyl, benzyl, pyridyl, piperidinyl, or cyclohexyl ring, wherein said ring is optionally substituted with benzyloxy, phenoxy, $-SO_2NH_2$, $-OH$, $-NO_2$, $-NH_2$, $-OMe$, $-Br$, $-Cl$, $-CO_2Me$, $-NHCO_2Me$, $-NHCO_2Et$, $-NHCON(Me)_2$, $-NHCON(Et)_2$, $-NHCOpyrrolidin-1-yl$, $-NHCOMorpholin-4-yl$, $-O-CH_2$ -phenyl, $-O(CH_2)_3OH$, $-O(CH_2)_3NH(CH_2)_2OH$, $-O(CH_2)_2NH(CH_2)_2OH$, $-O(CH_2)_3N(hydroxyethyl)(methyl)$, $-O(CH_2)_3pyrrolidin-1-yl$, $-O(CH_2)_2morpholin-4-yl$, $-O(CH_2)_3N(Me)_2$, $-O(CH_2)_3N(Et)_2$, $-O(CH_2)_3(4-hydroxyethyl piperazin-1-yl)$, $-O(CH_2)_3piperazin-1-yl$, $-O(CH_2)_3(4-hydroxymethylpiperidin-1-yl)$, $-O(CH_2)_3(4-hydroxypiperidin-1-yl)$, $-NHCO(CH_2)_3N(Me)_2$, $-NHCO(CH_2)_3NCOCH_3$, $-NHCOCH_2pyridin-2-yl$, $-NHCOCH_2(2-aminothiazol-4-yl)$, $-NHCOCH_2cyclopropyl$, $-NHCO(CH_2)_2N(Et)_2$, $-NHCO(CH_2)_2$ -(piperazin-2,5-dione-3-yl), $-NHCO_2CH_2tetrahydrofuran-2-yl$, $-NHCO_2tetrahydrofuran-2-yl$, $-NHCO_2tetrahydropyran-4-yl$, or $-NHCO_2CH_2tetrahydropyran-2-yl$;
- (b) R^2 is selected from phenyl, pyridyl, pyrimidinyl, cyclohexyl, piperidinyl, furanyl, or benzyl, wherein R^2 is optionally substituted with phenyl, phenoxy, benzyl, benzyloxy, pyridyl, 3-hydroxyphenyl, 2-hydroxyphenyl, 3-aminophenyl, N-BOC-pyrrolyl, 4-chlorophenyl, 3-ethoxypyridyl, 2-methoxypyridyl, 2,5-

dimethylisoxazolyl, 3-ethoxyphenyl, 4-isopropylphenyl, 4-F-3-Cl-phenyl, pyrrolyl, pyrimidinyl, chloro, bromo, fluoro, trifluoromethyl, -OH, -NH₂, methyl, methoxy, or ethoxy; and

(c) G is hydrogen.

5. (Original) The compound according to claim 1, wherein said compound has the formula **IIb**:



IIb

or a pharmaceutically acceptable derivative thereof.

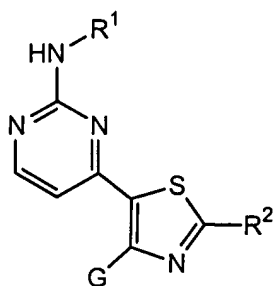
6. (Original) The compound according to claim 5, wherein said compound has one or more features selected from the group consisting of:

- (a) R¹ is hydrogen, Ar¹ or -T-Ar¹ wherein T is a C₁₋₄ alkylidene chain and Ar¹ is a 6-membered saturated, partially saturated, or aryl ring having zero to two heteroatoms independently selected from nitrogen, oxygen, or sulfur, and wherein each member of R¹ is optionally substituted with one -Z-R³ and one to three additional groups independently selected from -CO₂R, -OR, halogen, -NRSO₂R, -SO₂N(R)₂, -NRCON(R)₂, -NO₂, or -N(R)₂;
- (b) R² is Ar² or -CH₂-Ar² wherein Ar² is selected from 5-6 membered ring selected from carbocyclic, aryl, or a heterocyclyl or heteroaryl ring having one to two heteroatoms independently selected from nitrogen, oxygen or sulfur, and wherein Ar² is optionally substituted with one to five groups independently selected from -Z-R³, -R, halogen, -NO₂, -CN, -OR, -SR, -N(R)₂, -NRC(O)R, -NRC(O)N(R)₂, -NRCO₂R, -C(O)R, -CO₂R, -C(O)N(R)₂, -OC(O)N(R)₂, -S(O)R, -SO₂R, -SO₂N(R)₂, -N(R)SO₂R, -N(R)SO₂N(R)₂, -C(O)C(O)R, or -C(O)CH₂C(O)R; and
- (c) G is hydrogen.

7. (Original) The compound according to claim 6, wherein said compound has one or more features selected from the group consisting of:

- (a) R^1 is selected from a phenyl, benzyl, pyridyl, piperidinyl, or cyclohexyl ring, wherein said ring is optionally substituted with benzyloxy, phenoxy, $-SO_2NH_2$, $-OH$, $-NO_2$, $-NH_2$, $-OMe$, $-Br$, $-Cl$, $-CO_2Me$, $-NHCO_2Me$, $-NHCO_2Et$, $-NHCON(Me)_2$, $-NHCON(Et)_2$, $-NHCO$ pyrrolidin-1-yl, $-NHCO$ morpholin-4-yl, $-O-CH_2$ -phenyl, $-O(CH_2)_3OH$, $-O(CH_2)_3NH(CH_2)_2OH$, $-O(CH_2)_2NH(CH_2)_2OH$, $-O(CH_2)_3N$ (hydroxyethyl)(methyl), $-O(CH_2)_3$ pyrrolidin-1-yl, $-O(CH_2)_2$ morpholin-4-yl, $-O(CH_2)_3N(Me)_2$, $-O(CH_2)_3N(Et)_2$, $-O(CH_2)_3$ (4-hydroxyethyl piperazin-1-yl), $-O(CH_2)_3$ piperazin-1-yl, $-O(CH_2)_3$ (4-hydroxymethylpiperidin-1-yl), $-O(CH_2)_3$ (4-hydroxypiperidin-1-yl), $-NHCO(CH_2)_3N(Me)_2$, $-NHCO(CH_2)_3NCOCH_3$, $-NHCOCH_2$ pyridin-2-yl, $-NHCOCH_2$ (2-aminothiazol-4-yl), $-NHCOCH_2$ cyclopropyl, $-NHCO(CH_2)_2N(Et)_2$, $-NHCO(CH_2)_2$ (piperazin-2,5-dione-3-yl), $-NHCO_2CH_2$ tetrahydrofuran-2-yl, $-NHCO_2$ tetrahydrofuran-2-yl, $-NHCO_2$ tetrahydropyran-4-yl, or $-NHCO_2CH_2$ tetrahydropyran-2-yl;
- (b) R^2 is selected from phenyl, pyridyl, pyrimidinyl, cyclohexyl, piperidinyl, furanyl, or benzyl, wherein R^2 is optionally substituted with phenyl, phenoxy, benzyl, benzyloxy, pyridyl, 3-hydroxyphenyl, 2-hydroxyphenyl, 3-aminophenyl, N-BOC-pyrrolyl, 4-chlorophenyl, 3-ethoxypyridyl, 2-methoxypyridyl, 2,5-dimethylisoxazolyl, 3-ethoxyphenyl, 4-isopropylphenyl, 4-F-3-Cl-phenyl, pyrrolyl, pyrimidinyl, chloro, bromo, fluoro, trifluoromethyl, $-OH$, $-NH_2$, methyl, methoxy, or ethoxy; and
- (c) G is hydrogen.

8. (Original) The compound according to claim 1, wherein said compound has the formula **IIc**:

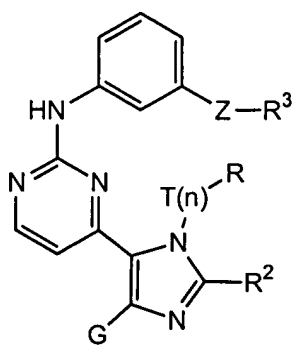


IIc

or a pharmaceutically acceptable derivative thereof.

9. to 11. (Canceled).

12. (Original) The compound according to claim 1, wherein said compound has the formula **IVa**:



IVa

or a pharmaceutically acceptable derivative thereof.

13. (Original) The compound according to claim 12, wherein said compound has one or more features selected from the group consisting of:

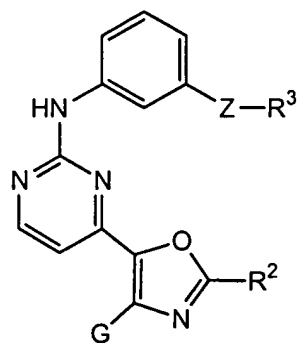
- (a) R^2 is Ar^2 or $-CH_2-Ar^2$ wherein Ar^2 is selected from 5-6 membered ring selected from carbocyclic, aryl, or a heterocyclyl or heteroaryl ring having one to two heteroatoms independently selected from nitrogen, oxygen or sulfur, and wherein Ar^2 is optionally substituted by wherein Ar^2 is optionally substituted with one to five groups independently selected from $-Z-R^3$, $-R$, halogen, $-NO_2$, $-CN$, $-OR$, $-SR$, $-N(R)_2$, $-NRC(O)R$, $-NRC(O)N(R)_2$, $-NRCO_2R$, $-C(O)R$, $-CO_2R$, $-C(O)N(R)_2$, $-OC(O)N(R)_2$, $-S(O)R$, $-SO_2R$, $-SO_2N(R)_2$, $-N(R)SO_2R$, $-N(R)SO_2N(R)_2$, $-C(O)C(O)R$, or $-C(O)CH_2C(O)R$;

- (b) G is hydrogen;
- (c) Z is a C₁₋₄ alkylidene chain wherein one methylene unit of Z is optionally replaced by -O-, -NH-, -NHC(O)-, -NHC(O)O-, -NHSO₂-, -C(O)NH-; and
- (d) R³ is selected from -N(R)₂, -NHC(O)R, or Ar³ wherein Ar³ is a 5-6 membered heterocyclic or heteroaryl ring having one to two heteroatoms independently selected from nitrogen, oxygen, or sulfur and Ar³ is optionally substituted with -R', -OR', -N(R')₂, or oxo.

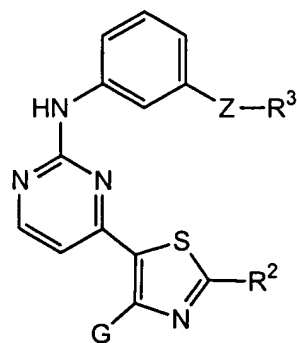
14. (Original) The compound according to claim 13, wherein said compound has one or more features selected from the group consisting of:

- (a) R² is selected from phenyl, pyridyl, pyrimidinyl, cyclohexyl, piperidinyl, furanyl, or benzyl, wherein each member of R² is optionally substituted with phenyl, phenoxy, benzyl, benzyloxy, pyridyl, 3-hydroxyphenyl, 2-hydroxyphenyl, 3-aminophenyl, N-BOC-pyrrolyl, 4-chlorophenyl, 3-ethoxypyridyl, 2-methoxypyridyl, 2,5-dimethylisoxazolyl, 3-ethoxyphenyl, 4-isopropylphenyl, 4-F-3-Cl-phenyl, pyrrolyl, pyrimidinyl, chloro, bromo, fluoro, trifluoromethyl, -OH, -NH₂, methyl, methoxy, or ethoxy;
- (b) G is hydrogen; and
- (c) -Z-R³ is selected from -O-CH₂-phenyl, -O(CH₂)₃OH, -O(CH₂)₃NH(CH₂)₂OH, -O(CH₂)₂NH(CH₂)₂OH, -O(CH₂)₃N(hydroxyethyl)(methyl), -O(CH₂)₃pyrrolidin-1-yl, -O(CH₂)₂morpholin-4-yl, -O(CH₂)₃N(Me)₂, -O(CH₂)₃N(Et)₂, -O(CH₂)₃(4-hydroxyethyl piperazin-1-yl), -O(CH₂)₃piperazin-1-yl, -O(CH₂)₃(4-hydroxymethylpiperidin-1-yl), -O(CH₂)₃(4-hydroxypiperidin-1-yl), -NHCO(CH₂)₃N(Me)₂, -NHCO(CH₂)₃NCOCH₃, -NHCOCH₂pyridin-2-yl, -NHCOCH₂(2-aminothiazol-4-yl), -NHCOCH₂cyclopropyl, -NHCO(CH₂)₂N(Et)₂, -NHCO(CH₂)₂-(piperazin-2,5-dione-3-yl), -NHC(O)-pyrrolidin-1-yl, -NHCOMorpholin-4-yl, -NHCO₂CH₂tetrahydrofuran-2-yl, -NHCO₂tetrahydrofuran-2-yl, -NHCO₂tetrahydropyran-4-yl, or -NHCO₂CH₂tetrahydropyran-2-yl.

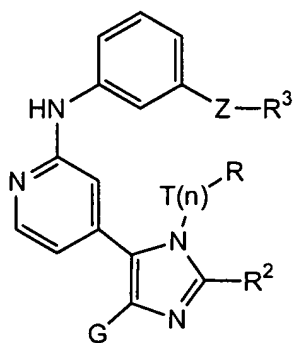
15. (Original) The compound according to claim 1, wherein said compound has the formula IVb, IVc, Va, Vb, or Vc:



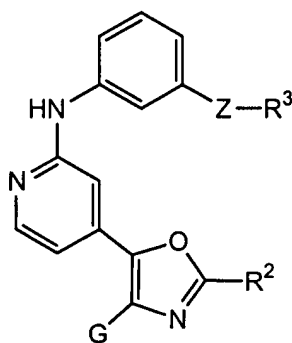
IVb



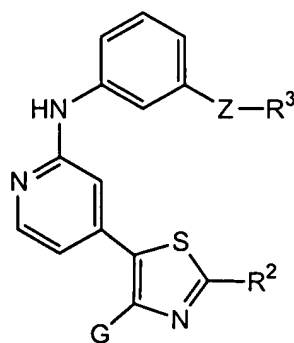
IVc



Va



Vb



Vc

or a pharmaceutically acceptable derivative thereof.

16. (Original) The compound according to claim 15, wherein said compound has one or more features selected from the group consisting of:

- (a) R^2 is Ar^2 or $-CH_2-Ar^2$ wherein Ar^2 is selected from 5-6 membered ring selected from carbocyclic, aryl, or a heterocyclyl or heteroaryl ring having one to two heteroatoms independently selected from nitrogen, oxygen or sulfur, and wherein Ar^2 is optionally substituted by wherein Ar^2 is optionally substituted with one to five groups independently selected from $-Z-R^3$, $-R$, halogen, $-NO_2$, $-CN$, $-OR$, $-SR$, $-N(R)_2$, $-NRC(O)R$, $-NRC(O)N(R)_2$, $-NRCO_2R$, $-C(O)R$, $-CO_2R$, $-C(O)N(R)_2$, $-OC(O)N(R)_2$, $-S(O)R$, $-SO_2R$, $-SO_2N(R)_2$, $-N(R)SO_2R$, $-N(R)SO_2N(R)_2$, $-C(O)C(O)R$, or $-C(O)CH_2C(O)R$;
- (b) G is hydrogen;
- (c) Z is a C_{1-4} alkylidene chain wherein one methylene unit of Z is optionally replaced by $-O-$, $-NH-$, $-NHC(O)-$, $-NHC(O)O-$, $-NHSO_2-$, $-C(O)NH-$; and

- (d) R^3 is selected from $-N(R)_2$, $-NHC(O)R$, or Ar^3 wherein Ar^3 is a 5-6 membered heterocyclic or heteroaryl ring having one to two heteroatoms independently selected from nitrogen, oxygen, or sulfur and Ar^3 is optionally substituted with $-R'$, $-OR'$, $-N(R')_2$, or oxo.
17. (Original) The compound according to claim 16, wherein said compound has one or more features selected from the group consisting of:
- (a) R^2 is selected from phenyl, pyridyl, pyrimidinyl, cyclohexyl, piperidinyl, furanyl, or benzyl, wherein each member of R^2 is optionally substituted with phenyl, phenoxy, benzyl, benzyloxy, pyridyl, 3-hydroxyphenyl, 2-hydroxyphenyl, 3-aminophenyl, N-BOC-pyrrolyl, 4-chlorophenyl, 3-ethoxypyridyl, 2-methoxypyridyl, 2,5-dimethylisoxazolyl, 3-ethoxyphenyl, 4-isopropylphenyl, 4-F-3-Cl-phenyl, pyrrolyl, pyrimidinyl, chloro, bromo, fluoro, trifluoromethyl, $-OH$, $-NH_2$, methyl, methoxy, or ethoxy;
- (b) G is hydrogen; and
- (c) $-Z-R^3$ is selected from $-O-CH_2$ -phenyl, $-O(CH_2)_3OH$, $-O(CH_2)_3NH(CH_2)_2OH$, $-O(CH_2)_2NH(CH_2)_2OH$, $-O(CH_2)_3N(hydroxyethyl)(methyl)$, $-O(CH_2)_3pyrrolidin-1-yl$, $-O(CH_2)_2morpholin-4-yl$, $-O(CH_2)_3N(Me)_2$, $-O(CH_2)_3N(Et)_2$, $-O(CH_2)_3(4-hydroxyethyl piperazin-1-yl)$, $-O(CH_2)_3piperazin-1-yl$, $-O(CH_2)_3(4-hydroxymethylpiperidin-1-yl)$, $-O(CH_2)_3(4-hydroxypiperidin-1-yl)$, $-NHCO(CH_2)_3N(Me)_2$, $-NHCO(CH_2)_3NCOCH_3$, $-NHCOCH_2pyridin-2-yl$, $-NHCOCH_2(2-aminothiazol-4-yl)$, $-NHCOCH_2cyclopropyl$, $-NHCO(CH_2)_2N(Et)_2$, $-NHCO(CH_2)_2-(piperazin-2,5-dione-3-yl)$, $-NHC(O)-pyrrolidin-1-yl$, $-NHCOMorpholin-4-yl$, $-NHCO_2CH_2tetrahydrofuran-2-yl$, $-NHCO_2tetrahydrofuran-2-yl$, $-NHCO_2tetrahydropyran-4-yl$, or $-NHCO_2CH_2tetrahydropyran-2-yl$.
18. (Currently amended) The compound according to claim 1 selected from those listed in Tables 1-3 [[1-5]].
19. (Original) A composition comprising a compound according to any one of claims 1 to 18, and a pharmaceutically acceptable carrier, adjuvant, or vehicle.

20. (Original) The composition according to claim 19, additionally comprising a therapeutic agent selected from an anti-proliferative agent, an anti-inflammatory agent, an immunomodulatory agent, a neurotrophic factor, an agent for treating cardiovascular disease, an agent for treating liver disease, an anti-viral agent, an agent for treating blood disorders, an agent for treating diabetes, an agent for treating immunodeficiency disorders, or an agent for treating cancer.

21. (Original) A method of inhibiting JNK, Lck, Src, or Aurora-2 kinase activity in a biological sample comprising the step of contacting said biological sample with:

- (a) a compound according to claim 1; or
- (b) a composition according to claim 19.

22. (Original) A method of treating or lessening the severity of a JNK-, Lck-, Src-, or Aurora-2-mediated disease or condition in a patient comprising the step of administering to said patient a composition according to claim 19.

23. (Original) A method of treating or lessening the severity of an inflammatory disease, autoimmune disease, destructive bone disorder, proliferative disorder, infectious disease, neurodegenerative disease, allergy, reperfusion/ischemia in stroke, heart attack, angiogenic disorder, organ hypoxia, vascular hyperplasia, cardiac hypertrophy, thrombin-induced platelet aggregation, or a condition associated with proinflammatory cytokines, comprising the step of administering to said patient a composition according to claim 19.

24. (Original) The method according to claim 23, wherein said method is used to treat or prevent an inflammatory disease selected from acute pancreatitis, chronic pancreatitis, asthma, allergies, or adult respiratory distress syndrome.

25. (Original) The method according to claim 23, wherein said method is used to treat or prevent an autoimmune disease selected from glomerulonephritis, rheumatoid arthritis, systemic lupus erythematosus, scleroderma, chronic thyroiditis, Graves' disease, autoimmune gastritis, diabetes, autoimmune hemolytic anemia, autoimmune neutropenia, thrombocytopenia, atopic dermatitis, chronic active hepatitis, myasthenia gravis, multiple sclerosis, inflammatory bowel disease, ulcerative colitis, Crohn's disease, psoriasis, or graft vs. host disease.

26. (Original) The method according to claim 23, wherein said method is used to treat or prevent a destructive bone disorders selected from osteoarthritis, osteoporosis or multiple myeloma-related bone disorder.

27. (Original) The method according to claim 23, wherein said method is used to treat or prevent a proliferative disease selected from acute myelogenous leukemia, chronic myelogenous leukemia, metastatic melanoma, Kaposi's sarcoma, multiple myeloma, .

28. (Original) The method according to claim 23, wherein said method is used to treat or prevent neurodegenerative disease selected from Alzheimer's disease, Parkinson's disease, amyotrophic lateral sclerosis, Huntington's disease, cerebral ischemia or neurodegenerative disease caused by traumatic injury, glutamate neurotoxicity or hypoxia.

29. (Original) The method according to claim 23, wherein said method is used to treat or prevent ischemia/reperfusion in stroke or myocardial ischemia, renal ischemia, heart attacks, organ hypoxia or thrombin-induced platelet aggregation.

30. (Original) The method according to claim 23, wherein said method is used to treat or prevent a condition associated with T-cell activation or pathologic immune responses.

31. (Original) The method according to claim 23, wherein said method is used to treat or prevent an angiogenic disorder selected from solid tumors, ocular neovascularization, or infantile haemangiomas.

32. (Original) The method according to claim 22, wherein said disease is selected from hypercalcemia, restenosis, hypercalcemia, osteoporosis, osteoarthritis, symptomatic treatment of bone metastasis, rheumatoid arthritis, inflammatory bowel disease, multiple sclerosis, psoriasis, lupus, graft vs. host disease, T-cell mediated hypersensitivity disease, Hashimoto's thyroiditis, Guillain-Barre syndrome, chronic obstructive pulmonary disorder, contact dermatitis, cancer, Paget's disease, asthma, ischemic or reperfusion injury, allergic disease, atopic dermatitis, or allergic rhinitis.

33. (Original) The method according to claim 32, wherein said disease is selected from hypercalcemia, osteoporosis, osteoarthritis, or symptomatic treatment of bone metastasis.

34. (Original) The method according to claim 22, wherein said disease is selected from autoimmune diseases, allergies, rheumatoid arthritis, or leukemia.

35. (Original) The method according to claim 22, wherein said disease is selected from melanoma, leukemia, or a cancer selected from colon, breast, gastric, ovarian, cervical, melanoma, renal, prostate, lymphoma, neuroblastoma, pancreatic, leukemia and bladder.

36. (Original) The method according to claim 22, comprising the additional step of administering to said patient an additional therapeutic agent selected from an anti-proliferative agent, an anti-inflammatory agent, an immunomodulatory agent, a neurotrophic factor, an agent for treating cardiovascular disease, an agent for treating liver disease, an anti-viral agent, an agent for treating blood disorders, an agent for treating diabetes, or an agent for treating immunodeficiency disorders, wherein:

said additional therapeutic agent is appropriate for the disease being treated; and

said additional therapeutic agent is administered together with said composition as a single dosage form or separately from said composition as part of a multiple dosage form.

37. (Original) A composition for coating an implantable device comprising a compound according to claim 1 and a carrier suitable for coating said implantable device.

38. (Canceled).